CLAIMS

1. Use of natural human α-interferon for the preparation of a medicament in liquid form to be administered through peroral route at dosages comprised between 100 UI and 500 UI/day, for therapy of viral hepatitis in humans and animals.

2. Use of natural human α-interferon for the preparation of a medicament in liquid form to be administered through peroral route at dosages comprised between 100 UI and 500 UI/day, for therapy of neoplasia and immunologic diseases in humans and animals.

3. Use of natural human α-interferon according to claims 1 or 2 wherein said interferon is obtained from lymphoblastoid cell cultures.

4. Use of natural human α-interferon according to claims 1 or 2 wherein said interferon is obtained from lymphocyte cells.

5. Use of natural human α-interferon according to any of previous claims wherein said medicament is administered in mono dosage units of appr. 1 ml.

6. Pharmaceutical liquid composition for peroral administration comprising natural human α-interferon either from lymphoblastoid cell cultures or from lymphocyte cells at a concentration between 100 UI/ml and 500 UI/ml.

PHARMACEUTICAL COMPOSITIONS COMPRISING NATURAL HUMAN α-INTERFERON

Use of natural human α-interferon for the preparation of a medicament in liquid form to be administered through peroral route at dosages comprised between 100 UI and 500 UI/day, for therapy of viral infections, in particular viral hepatitis, neoplasia and immune diseases in humans and animals.